L26

L27

L28

30 FILE CAPLUS

6 FILE PCTFULL

2 FILE EUROPATFULL

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TOTAL FOR ALL FILES
      38 S E29 OR E31-33
L30
             3 FILE CAPLUS
L31
             O FILE EUROPATFULL
L32
             1 FILE PCTFULL
    TOTAL FOR ALL FILES
            4 S L29 AND ADRENERGIC
L33
L34
          1418 FILE CAPLUS
L35
            27 FILE EUROPATFULL
L36
            66 FILE PCTFULL
    TOTAL FOR ALL FILES
          1511 S METHOXAMINE AND ADRENERGIC
L37
          1011 FILE CAPLUS
L38
           26 FILE EUROPATFULL
L39
L40
            64 FILE PCTFULL
    TOTAL FOR ALL FILES
          1101 S L37 AND (ADRENERGIC (2A) ANTAGON? OR BLOC? OR INHIBIT?)
L41
     FILE 'CAPLUS' ENTERED AT 20:02:15 ON 30 MAY 2002
          152 S METHOXAMINE AND (ADRENERGIC ANTAGON?)
L42
            66 S METHOXAMINE (1S) (ADRENERGIC ANTAGON?)
L43
    FILE 'JAPIO' ENTERED AT 20:15:31 ON 30 MAY 2002
          33 S PHENTOLAMINE OR PRAZOSIN OR DOXAZOSIN
L44
        252516 S (ADRENERGIC (2A) ANTAGON? OR BLOC? OR INHIBIT?)
L45
            82 S (ADRENERGIC (2A) (ANTAGON? OR BLOC? OR INHIBIT?))
L46
L47
           114 S L44 OR L46
             0 S L47 AND L7
L48
     FILE 'MEDLINE, SCISEARCH, BIOSIS, EMBASE' ENTERED AT 20:17:24 ON 30 MAY
     2002
L49
         55921 FILE MEDLINE
L50
         16840 FILE SCISEARCH
L51
         38167 FILE BIOSIS
L52
         82563 FILE EMBASE
    TOTAL FOR ALL FILES
    193491 S L47
L53
L54
            78 FILE MEDLINE
L55
            33 FILE SCISEARCH
L56
            65 FILE BIOSIS
            95 FILE EMBASE
L57
    TOTAL FOR ALL FILES
    271 S L47 AND L7
L58
            6 FILE MEDLINE
L59
L60
             6 FILE SCISEARCH
L61
             3 FILE BIOSIS
L62
             9 FILE EMBASE
    TOTAL FOR ALL FILES
L63
            24 S L58 NOT ANAL.
    FILE 'MEDLINE, SCISEARCH, BIOSIS, EMBASE, CAPLUS, USPATFULL' ENTERED AT
     20:26:27 ON 30 MAY 2002
         24368 FILE MEDLINE
L64
L65
          9983 FILE SCISEARCH
L66
         24349 FILE BIOSIS
L67
         42094 FILE EMBASE
L68
         20798 FILE CAPLUS
L69
          2578 FILE USPATFULL
    TOTAL FOR ALL FILES
     124170 S PHENTOLAMINE OR PRAZOSIN OR DOXAZOSIN OR ERGOTAMINE OR DIHYDR
L70
       24696 FILE MEDLINE
L71
L72
         8366 FILE SCISEARCH
    14366 FILE BIOSIS
L73
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L74
        80202 FILE EMBASE
       14296 FILE CAPLUS
L75
        39532 FILE USPATFULL
L76
    TOTAL FOR ALL FILES
    181458 S (RECTAL OR TOPICAL) (10A) (APPL? OR ADMINIST?)
L77
            79 FILE MEDLINE
L78
            28 FILE SCISEARCH
L79
            70 FILE BIOSIS
L80
L81
            73 FILE EMBASE
            48 FILE CAPLUS
L82
            36 FILE USPATFULL
L83
    TOTAL FOR ALL FILES
     334 S L70 (1S) L77
L84
           156 DUP REM L84 (178 DUPLICATES REMOVED)
L85
           79 S L85
L86
             3 FILE MEDLINE
L87
            5 S L85
L88
            0 FILE SCISEARCH
L89
L90
            8 S L85
            O FILE BIOSIS
L91
L92
           11 S L85
L93
             O FILE EMBASE
L94
           19 S L85
             O FILE CAPLUS
L95
            34 S L85
L96
             5 FILE USPATFULL
L97
    TOTAL FOR ALL FILES
L98
            8 S L85 AND L7
\Rightarrow s 198 not anal.
            O FILE MEDLINE
L99
            O FILE SCISEARCH
L100
L101
            O FILE BIOSIS
            O FILE EMBASE
L102
L103
            O FILE CAPLUS
L104
            O FILE USPATFULL
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TOTAL FOR ALL FILES

L105 0 L98 NOT ANAL.

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ANSWER 1 OF 1 REGISTRY COPYRIGHT 2002 ACS
L16
     50-60-2 REGISTRY
RN
     Phenol, 3-[[(4,5-dihydro-1H-imidazol-2-yl)methyl](4-methylphenyl)amino]-
CN
     (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
     Phenol, m-[N-(2-imidazolin-2-ylmethyl)-p-toluidino]- (8CI)
OTHER NAMES:
     2-(m-Hydroxy-N-p-tolylanilinomethyl)-2-imidazoline
CN
     2-(N'-p-Tolyl-N'-m-hydroxyphenylaminomethyl)-2-imidazoline
CN
     2-[[N-(m-Hydroxyphenyl)-p-toluidino]methyl]-2-imidazoline
CN
     C 7337
CN
     C 7337 Ciba
CN
     Dibasin
CN
     Fentolamine
CN
CN
     Phentolamine
CN
     Regitin
CN
     Regitine
FS
     3D CONCORD
MF
     C17 H19 N3 O
     COM
CI
                  ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*,
LC
     STN Files:
       BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CBNB, CEN, CHEMCATS, CHEMLIST, CIN, CSCHEM, DDFU, DIOGENES, DRUGNL,
       DRUGPAT, DRUGU, DRUGUPDATES, EMBASE, HSDB*, IFICDB, IFIUDB, IPA,
       MEDLINE, MRCK*, NIOSHTIC, PROMT, RTECS*, SYNTHLINE, TOXCENTER, USAN,
       USPATFULL, VETU
          (*File contains numerically searchable property data)
                     EINECS**, WHO
     Other Sources:
          (**Enter CHEMLIST File for up-to-date regulatory information)
```

$$CH_2-N$$
OH
Me

=>

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2402 REFERENCES IN FILE CA (1967 TO DATE)
3 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
2403 REFERENCES IN FILE CAPLUS (1967 TO DATE)
48 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

L5 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2002 ACS

RN 74191-85-8 REGISTRY

CN Piperazine, 1-(4-amino-6,7-dimethoxy-2-quinazolinyl)-4-[(2,3-dihydro-1,4-benzodioxin-2-yl)carbonyl]- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN 1,4-Benzodioxin, piperazine deriv.

OTHER NAMES:

CN (.+-.)-Doxazosin

CN Doxazosin

CN UK 33,274

CN UK 33274

FS 3D CONCORD

DR 137888-77-8

MF C23 H25 N5 O5

CI COM

=>

LC STN Files: ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*,
BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB,
CEN, CHEMCATS, CIN, CSCHEM, DDFU, DIOGENES, DRUGPAT, DRUGU, DRUGUPDATES,
EMBASE, IPA, MEDLINE, MRCK*, PHAR, PROMT, SYNTHLINE, TOXCENTER, USAN,
USPATFULL, VETU

(*File contains numerically searchable property data) Other Sources: WHO

$$\begin{array}{c|c} & & & & \\ & & & \\ \text{MeO} & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & \\ & & & \\ & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & &$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

404 REFERENCES IN FILE CA (1967 TO DATE)

5 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

406 REFERENCES IN FILE CAPLUS (1967 TO DATE)

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L3
     ANSWER 1 OF 1 REGISTRY COPYRIGHT 2002 ACS
     19216-56-9 REGISTRY
RN
     Piperazine, 1-(4-amino-6,7-dimethoxy-2-quinazolinyl)-4-(2-furanylcarbonyl)-
CN
      (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
     Piperazine, 1-(4-amino-6,7-dimethoxy-2-quinazolinyl)-4-(2-furoyl)- (8CI)
OTHER NAMES:
     Lentopres
CN
     Prazosin
CN
     3D CONCORD
FS
     C19 H21 N5 O4
MF
CI
     COM
                  ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS,
LC
     STN Files:
       BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB, CEN, CHEMCATS,
       CHEMLIST, CIN, CSCHEM, DDFU, DIOGENES, DRUGNL, DRUGPAT, DRUGU,
       DRUGUPDATES, EMBASE, HSDB*, IFICDB, IFIPAT, IFIUDB, IPA, MEDLINE, MRCK*,
       NAPRALERT, NIOSHTIC, PHAR, PROMT, RTECS*, SPECINFO, TOXCENTER, USAN,
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(*File contains numerically searchable property data)
Other Sources: EINECS**, WHO

(**Enter CHEMLIST File for up-to-date regulatory information)

USPATFULL, VETU

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1938

L5 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS 2000:383138 CAPLUS 134:61209 DN ΤI An innovative cosmeceutical with skin whitening activity. Note ΑΠ Morganti, P.; Fabrizi, G.; James, B. President/Director, R. and D - Mavi Sud S.r.l., Aprilia, 04011, Italy CS Journal of Applied Cosmetology (1999), 17(4), 144-153 CODEN: JACOEL; ISSN: 0392-8543 PB International Ediemme Journal DT LA English CC 62-4 (Essential Oils and Cosmetics) Section cross-reference(s): 63 AB Hyperpigmentation is a skin disturbance affecting many people all over the world. Among the different bleaching cosmetic products, the most commonly used active ingredients are hydroquinone, azelaic acid, kojic acid, ellagic acid, rucinol, arbutin and different vitamin C derivs. In fact, vitamin C is widely known to have a suppressing effect on melanic pigmentation, but because of its easy decompn., a variety of stabilized vitamin C derivs. have been developed and commercialized. The main problem of these derivs. is their difficulty to target the stratum corneum (SC) for acting specifically on functioning melanocytes with active synthesis of melanin. The aim of this study was to control the combined activity of arbutin ext., hexadecanoyl ascorbic acid (VC-IP) and magnesium L-ascorbyl-2-phosphate (VC-PMG), to suppress melanic pigmentation (product A). At the same time, we wanted to control the depigmenting activity and the product stability of the ascorbic-acid, included in a kojic-based cosmetic formulation utilizing a new 2-chamber dispenser (SYMBIO), which allows to keep vitamin C sep. from the other ingredients (product B). Skin absorption-potential through the skin of the cosmetic vehicles and active ingredients were controlled by the dansyl chloride methodol., stripping the SC at different levels. Clin. evaluation of the obtained lightening effect was performed on 40 randomized female volunteers over a period of 3 mo by the clin. score and the Minolta Chromameter CR 200 methods. The topical application of both the products (A and B) was effective in lightening the skin of the majority of the treated patients, showing a remarkable penetrability degree and a mean redn. of the skin hyperpigmentation from 30 to 45%. L-ascorbic acid-based formulation was superior of about 20% to VC-PMG-based in restoring to normal the hyperpigmentation skin disorders, such as senile freckles. Both the formulations were well tolerated during the study term. ST ascorbate cosmeceutical skin lightening; palmitate ascorbate cosmeceutical skin ΙT Skin, disease (hyperpigmentation; cosmeceutical with skin lightening activity) Cosmetics (skin-lightening; cosmeceutical with skin lightening activity) IT Skin (stratum corneum; cosmeceutical with skin lightening activity) IT Drug delivery systems (topical; cosmeceutical with skin lightening activity) 50-81-7, L-Ascorbic acid, biological studies 137-66-6, Ascorbyl 23666-04-8, Magnesium ascorbyl-2-phosphate RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BUU (Biological use, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (cosmeceutical with skin lightening activity) RE.CNT THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD 19 RE (1) Bose, S; Cosmetic Dermatology 1994, P277 (2) Colton, T; Statistics in medicine 1974

- (3) Darr, D; Br J Dermatol 1992, V127, P247 CAPLUS
- (4) Edens, L; J Appl Cosmetol 1999, V17, P1
- (5) Funasaka, Y; Fragrance J 1997, V1997-9, P19

- (6) Kameyama, K; J Am Acad Dermatol 1996, V34, P29 MEDLINE (7) Koichiro, K; J Am Acad Dermatol 1996, V34, P29 (8) Maeda, K; J Pharmacol Exp Therap 1996, V276, P765 CAPLUS
- (9) Matoba, M; Proceedings 4th Scientific Conference of the Asian Societies of Cosmetic Scientists 1999, P136
- (10) Mc Callagh, P; J R Stat Soc Ser B 1980, V42, P109
- (11) Mishima, Y; Skin Research 1994, V36, P134 CAPLUS
- (12) Morganti, P; Cosmet & Toilet 1997, V112, P61 CAPLUS (13) Morganti, P; J Appl Cosmetol 1997, V15, P147 CAPLUS (14) Okubo, T; J Dermatological Science 1995, V10, P88

- (15) Ortonne, J; Aesthetic Dermatology 1991, P74
- (16) Perricone, N; J Geriatric Derm 1997, V5(4), P162
- (17) Ridge, B; Br J Dermatol 1988, V118, P167 CAPLUS
- (18) Shinomiya, T; Fragrance J 1997, V1997-3, P80 (19) Tachibana, S; Fragrance J 1997, V1997-9, P37

ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS L_5 2000:383138 CAPLUS 134:61209 DN ΤI An innovative cosmeceutical with skin whitening activity. Note ΑU Morganti, P.; Fabrizi, G.; James, B. President/Director, R. and D - Mavi Sud S.r.l., Aprilia, 04011, Italy CS Journal of Applied Cosmetology (1999), 17(4), 144-153 SO CODEN: JACOEL; ISSN: 0392-8543 PR International Ediemme DT Journal English LA 62-4 (Essential Oils and Cosmetics) CC Section cross-reference(s): 63 Hyperpigmentation is a skin disturbance affecting many people all over the AB world. Among the different bleaching cosmetic products, the most commonly used active ingredients are hydroquinone, azelaic acid, kojic acid, ellagic acid, rucinol, arbutin and different vitamin C derivs. In fact, vitamin C is widely known to have a suppressing effect on melanic pigmentation, but because of its easy decompn., a variety of stabilized vitamin C derivs. have been developed and commercialized. The main problem of these derivs. is their difficulty to target the stratum corneum (SC) for acting specifically on functioning melanocytes with active synthesis of melanin. The aim of this study was to control the combined activity of arbutin ext., hexadecanoyl ascorbic acid (VC-IP) and magnesium L-ascorbyl-2-phosphate (VC-PMG), to suppress melanic pigmentation (product A). At the same time, we wanted to control the depigmenting activity and the product stability of the ascorbic-acid, included in a kojic-based cosmetic formulation utilizing a new 2-chamber dispenser (SYMBIO), which allows to keep vitamin C sep. from the other ingredients (product B). Skin absorption-potential through the skin of the cosmetic vehicles and active ingredients were controlled by the dansyl chloride methodol., stripping the SC at different levels. Clin. evaluation of the obtained lightening effect was performed on 40 randomized female volunteers over a period of 3 mo by the clin. score and the Minolta Chromameter CR 200 methods. The topical application of both the products (A and B) was effective in lightening the skin of the majority of the treated patients, showing a remarkable penetrability degree and a mean redn. of the skin hyperpigmentation from 30 to 45%. L-ascorbic acid-based formulation was superior of about 20% to VC-PMG-based in restoring to normal the hyperpigmentation skin disorders, such as senile freckles. Both the formulations were well tolerated during the study term. ST ascorbate cosmeceutical skin lightening; palmitate ascorbate cosmeceutical skin ΙT Skin, disease (hyperpigmentation; cosmeceutical with skin lightening activity) TΤ Cosmetics (skin-lightening; cosmeceutical with skin lightening activity) TT (stratum corneum; cosmeceutical with skin lightening activity) Drug delivery systems (topical; cosmeceutical with skin lightening activity) 50-81-7, L-Ascorbic acid, biological studies 137-66-6, Ascorbyl TT palmitate 23666-04-8, Magnesium ascorbyl-2-phosphate RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BUU (Biological use, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (cosmeceutical with skin lightening activity) THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 19

(1) Bose, S; Cosmetic Dermatology 1994, P277(2) Colton, T; Statistics in medicine 1974

- (3) Darr, D; Br J Dermatol 1992, V127, P247 CAPLUS
- (4) Edens, L; J Appl Cosmetol 1999, V17, Pl
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- (9) Matoba, M; Proceedings 4th Scientific Conference of the Asian Societies of Cosmetic Scientists 1999, P136
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- (11) Mishima, Y; Skin Research 1994, V36, P134 CAPLUS
- (12) Morganti, P; Cosmet & Toilet 1997, V112, P61 CAPLUS (13) Morganti, P; J Appl Cosmetol 1997, V15, P147 CAPLUS (14) Okubo, T; J Dermatological Science 1995, V10, P88
- (15) Ortonne, J; Aesthetic Dermatology 1991, P74
- (16) Perricone, N; J Geriatric Derm 1997, V5(4), P162
- (17) Ridge, B; Br J Dermatol 1988, V118, P167 CAPLUS
- (18) Shinomiya, T; Fragrance J 1997, V1997-3, P80 (19) Tachibana, S; Fragrance J 1997, V1997-9, P37

 L_5 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS AN 2000:383138 CAPLUS DN 134:61209 TI An innovative cosmeceutical with skin whitening activity. Note ΑU Morganti, P.; Fabrizi, G.; James, B. President/Director, R. and D - Mavi Sud S.r.l., Aprilia, 04011, Italy CS Journal of Applied Cosmetology (1999), 17(4), 144-153 SO CODEN: JACOEL; ISSN: 0392-8543 International Ediemme PR DT Journal LA English 62-4 (Essential Oils and Cosmetics) CC Section cross-reference(s): 63 Hyperpigmentation is a skin disturbance affecting many people all over the AB world. Among the different bleaching cosmetic products, the most commonly used active ingredients are hydroquinone, azelaic acid, kojic acid, ellagic acid, rucinol, arbutin and different vitamin C derivs. In fact, vitamin C is widely known to have a suppressing effect on melanic pigmentation, but because of its easy decompn., a variety of stabilized vitamin C derivs. have been developed and commercialized. The main problem of these derivs. is their difficulty to target the stratum corneum (SC) for acting specifically on functioning melanocytes with active synthesis of melanin. The aim of this study was to control the combined activity of arbutin ext., hexadecanoyl ascorbic acid (VC-IP) and magnesium L-ascorbyl-2-phosphate (VC-PMG), to suppress melanic pigmentation (product A). At the same time, we wanted to control the depigmenting activity and the product stability of the ascorbic-acid, included in a kojic-based cosmetic formulation utilizing a new 2-chamber dispenser (SYMBIO), which allows to keep vitamin C sep. from the other ingredients (product B). Skin absorption-potential through the skin of the cosmetic vehicles and active ingredients were controlled by the dansyl chloride methodol., stripping the SC at different levels. Clin. evaluation of the obtained lightening effect was performed on 40 randomized female volunteers over a period of 3 mo by the clin. score and the Minolta Chromameter CR 200methods. The topical application of both the products (A and B) was effective in lightening the skin of the majority of the treated patients, showing a remarkable penetrability degree and a mean redn. of the skin hyperpigmentation from 30 to 45%. L-ascorbic acid-based formulation was superior of about 20% to VC-PMG-based in restoring to normal the hyperpigmentation skin disorders, such as senile freckles. Both the formulations were well tolerated during the study term. ascorbate cosmeceutical skin lightening; palmitate ascorbate cosmeceutical ST skin ΤT Skin, disease (hyperpigmentation; cosmeceutical with skin lightening activity) ΙT Cosmetics (skin-lightening; cosmeceutical with skin lightening activity) ΙT Skin (stratum corneum; cosmeceutical with skin lightening activity) Drug delivery systems ΙT (topical; cosmeceutical with skin lightening activity) IT 50-81-7, L-Ascorbic acid, biological studies 137-66-6, Ascorbyl 23666-04-8, Magnesium ascorbyl-2-phosphate RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BUU (Biological use, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (cosmeceutical with skin lightening activity) RE.CNT THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD RE (1) Bose, S; Cosmetic Dermatology 1994, P277

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- (5) Funasaka, Y; Fragrance J 1997, V1997-9, P19(6) Kameyama, K; J Am Acad Dermatol 1996, V34, P29 MEDLINE
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- (8) Maeda, K; J Pharmacol Exp Therap 1996, V276, P765 CAPLUS
- (9) Matoba, M; Proceedings 4th Scientific Conference of the Asian Societies of Cosmetic Scientists 1999, P136

- (10) Mc Callagh, P; J R Stat Soc Ser B 1980, V42, P109 (11) Mishima, Y; Skin Research 1994, V36, P134 CAPLUS (12) Morganti, P; Cosmet & Toilet 1997, V112, P61 CAPLUS
- (13) Morganti, P; J Appl Cosmetol 1997, V15, P147 CAPLUS
- (14) Okubo, T; J Dermatological Science 1995, V10, P88
- (15) Ortonne, J; Aesthetic Dermatology 1991, P74
- (16) Perricone, N; J Geriatric Derm 1997, V5(4), P162 (17) Ridge, B; Br J Dermatol 1988, V118, P167 CAPLUS (18) Shinomiya, T; Fragrance J 1997, V1997-3, P80 (19) Tachibana, S; Fragrance J 1997, V1997-9, P37

L5 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS 2000:383138 CAPLUS ΑN 134:61209 DN An innovative cosmeceutical with skin whitening activity. Note TΙ Morganti, P.; Fabrizi, G.; James, B. ΑU President/Director, R. and D - Mavi Sud S.r.l., Aprilia, 04011, Italy CS Journal of Applied Cosmetology (1999), 17(4), 144-153 SO CODEN: JACOEL; ISSN: 0392-8543 PΒ International Ediemme DT Journal LAEnglish 62-4 (Essential Oils and Cosmetics) CC Section cross-reference(s): 63 Hyperpigmentation is a skin disturbance affecting many people all over the AΒ world. Among the different bleaching cosmetic products, the most commonly used active ingredients are hydroquinone, azelaic acid, kojic acid, ellagic acid, rucinol, arbutin and different vitamin C derivs. In fact, vitamin C is widely known to have a suppressing effect on melanic pigmentation, but because of its easy decompn., a variety of stabilized vitamin C derivs. have been developed and commercialized. The main problem of these derivs. is their difficulty to target the stratum corneum (SC) for acting specifically on functioning melanocytes with active synthesis of melanin. The aim of this study was to control the combined activity of arbutin ext., hexadecanoyl ascorbic acid (VC-IP) and magnesium L-ascorbyl-2-phosphate (VC-PMG), to suppress melanic pigmentation (product A). At the same time, we wanted to control the depigmenting activity and the product stability of the ascorbic-acid, included in a kojic-based cosmetic formulation utilizing a new 2-chamber dispenser (SYMBIO), which allows to keep vitamin C sep. from the other ingredients (product B). Skin absorption-potential through the skin of the cosmetic vehicles and active ingredients were controlled by the dansyl chloride methodol., stripping the SC at different levels. Clin. evaluation of the obtained lightening effect was performed on 40 randomized female volunteers over a period of 3 mo by the clin. score and the Minolta Chromameter CR 200 methods. The topical application of both the products (A and B) was effective in lightening the skin of the majority of the treated patients, showing a remarkable penetrability degree and a mean redn. of the skin hyperpigmentation from 30 to 45%. L-ascorbic acid-based formulation was superior of about 20% to VC-PMG-based in restoring to normal the hyperpigmentation skin disorders, such as senile freckles. Both the formulations were well tolerated during the study term. ST ascorbate cosmeceutical skin lightening; palmitate ascorbate cosmeceutical skin ΙT Skin, disease (hyperpigmentation; cosmeceutical with skin lightening activity) ΙT (skin-lightening; cosmeceutical with skin lightening activity) IT (stratum corneum; cosmeceutical with skin lightening activity) IT Drug delivery systems (topical; cosmeceutical with skin lightening activity) 50-81-7, L-Ascorbic acid, biological studies 137-66-6, Ascorbyl IT 23666-04-8, Magnesium ascorbyl-2-phosphate palmitate RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BUU (Biological use, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (cosmeceutical with skin lightening activity) RE.CNT THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD RE (1) Bose, S; Cosmetic Dermatology 1994, P277

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- (4) Edens, L; J Appl Cosmetol 1999, V17, P1
- (5) Funasaka, Y; Fragrance J 1997, V1997-9, P19
- (6) Kameyama, K; J Am Acad Dermatol 1996, V34, P29 MEDLINE
- (7) Koichiro, K; J Am Acad Dermatol 1996, V34, P29
- (8) Maeda, K; J Pharmacol Exp Therap 1996, V276, P765 CAPLUS
- (9) Matoba, M; Proceedings 4th Scientific Conference of the Asian Societies of Cosmetic Scientists 1999, P136
- (10) Mc Callagh, P; J R Stat Soc Ser B 1980, V42, P109
- (11) Mishima, Y; Skin Research 1994, V36, P134 CAPLUS
- (12) Morganti, P; Cosmet & Toilet 1997, V112, P61 CAPLUS
- (13) Morganti, P; J Appl Cosmetol 1997, V15, P147 CAPLUS
- (14) Okubo, T; J Dermatological Science 1995, V10, P88
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